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WPI Acc no: 1997-145233/199713 XRAM Acc no: C1997-046296

4,5-Diaryloxazol-2-yl- alkyl-aryl or -aryloxy alkanoic acid derivs. - are platelet

aggregation inhibitors, vasodilators, and antihypertensives Patent Assignee: FUJISAWA PHARM CO LTD (FUJI)

Inventor: HATTORI K; OKITSU O; TABUCHI S; TANIGUCHI K; TANIQUCHI K; TSUBAKI K
Patent Family (19 patents, 28 countries)

Patent Number   Kind   Date   Application Number   Kind   Date   Update   Typ   WO 1997003973   A1   19970206   WO 1996JP1996   A   19960718   199713   B   AU 199664697   A   19970218   AU 199664697   A   19960718   199723   E   ZA 199606126   A   19970430   ZA 19966126   A   19960718   199723   E   EP 842161   A1   19980520   EP 1996924137   A   19960718   199824   E   WO 1996JP1996   A   19960718   199824   E   WO 1996JP1996   A   19960718   1999081   A   19990881   A   19990881   A   19990881   A   19990818   A   19990718   I99940   E   HU 1999881   A   19960718   I99940   E   JP 11509191   W   19990817   WO 1996JP1996   A   19960718   I99943   E   JP 1997504319   A   19960718   I99952   E   US 1998983139   A   19980718   I99952   E   US 1998983139   A   19980120   Z00004   E   KR 1999063582   A   19990726   WO 1996JP1996   A   19960718   Z00020   E   KR 1999063582   A   19990726   WO 1996JP1996   A   19960718   Z00020   E   KR 19980344   B1   Z0011009   US 1998983139   A   19980120   Z00162   E   US 1999357664   A   19980717   Z00116   E   US 1999357664   A   19980717   Z00116   E   US 1999357664   A   19980718   Z00239   E   EP 20023081   A   19980718   Z00239   E   EP 20023081   A   19980718   Z00269   E   EP 842161   B1   Z0020918   EP 1996924137   A   19980718   Z00269   E   WO 1996JP1996   A   19980718   Z00269   E   EP 842161   B1   Z0020918   EP 1996924137   A   19980718   Z00269   E   EP 842161   B1   Z0020918   EP 1996924137   A   19980718   Z00269   E   EP 842161   B1   Z0020918   EP 1996924137   A   19980718   Z00269   E   EP 842161   B1   Z0020918   EP 1996924137   A   19980718   Z00269   E   EP 842161   B1   Z0020918   EP 1996924137   A   19980718   Z00269   E   WO 1996JP1996   A   19980718   Z00269   E   Z0026918   Z00269		Pa	atent Famil	ly (19 paterits, 26 Co	Junta	es /		
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Priority Applications (no., kind, date): WO 1996JP1996 A 19960718; GB 199515085 A 19950721; AU 19969002 A 19960329

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National Designated States,Original	ates,Original					
Regional Designated States Original	AT E	BE CI	H DE	DK E	A ES FI FR GB GR IE IT I	
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Regional Designated States, Original	AT	BE C	H D	E DK I	ES FI FR GB GR IE IT LI L	_U NL PT SE
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		PCT Application	WO 1996JP1996	
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### Alerting Abstract WO A1

(4,5-Diaryloxazol-2-ylalkyl)aryl (or -aryloxy) alkanoic acid derivs. of formula (I) and their sa protected); R2, R3 = aryl (opt. substd.); R4 = H, OH, 1-6C alkyl, or aryl; A1 = 1-6C alkylene substd. by R4); A3 = A4-A5; A4 = a bond, CH2, or CO; A5 = 5-8C cycloalkene, 7-8C cycloalkene, or piperidine (all opt. substd.), or X-A6; X = O, S, or NR5; R5 = H, 1-6C alkyl, or USE - (I) have platelet aggregation, vasodilating, and antihypertensive activities, and are protections, arterial obstruction, cerebrovascular disorders, gastric ulcer, hepatitis, hepatic restenosis after PTCA, hypertension, inflammation, autoimmune disease, heart failure, rena bedsores, and for protecting organs after transplant or surgery, also in organ preserving flu

Title Terms /Index Terms/Additional Words: DI; YL; ALKYL; ARYL; ARYLOXY; ALKANOIC; ANTIHYPERTENSIVE

### Class Codes

### International Patent Classification

IPC	Class Level	Scope	Position	Status	Version Date
A61K-031/33; A61K-031/42; C07D-263/32		·	Main	1.01	"Version 7"
A61K-031/00; C07D-263/34; C07D-413/04			Secondary		"Version 7"
A61K-0031/42	Α	I e		R	20060101
C07D-0263/32	Α	]		R	20060101
C07D-0413/04	Α	I		R	20060101
A61K-0031/42	С	1		R	20060101
C07D	S	1		R	20060101
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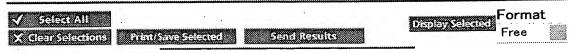
US Classification, Issued: 514326000, 514374000, 546209000, 548235000, 514326000, 5143

File Segment: CPI DWPI Class: B03

Manual Codes (CPI/A-N): B07-E01; B14-C03; B14-E08; B14-F01E; B14-F02; B14-F02D; I

R02; B14-S04

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# WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6:

C07D 263/32, A61K 31/42, C07D 413/04

(11) International Publication Number:

WO 97/03973

A1

(43) International Publication Date:

6 February 1997 (06.02.97)

(21) International Application Number:

PCT/JP96/01996

(22) International Filing Date:

.18 July 1996 (18.07.96)

(81) Designated States: AU, CA, CN, HU, JP, KR, MX, US, Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TI, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB,

GR, IE, IT, LU, MC, NL, PT, SE).

(30) Priority Data:

9515085.0 PN 9002 21 July 1995 (21.07.95) 29 March 1996 (29.03.96) GB AU

Published

With international search report.

(71) Applicant (for all designated States except US): FUJISAWA PHARMACEUTICAL CO., LTD. [JP/JP]; 4-7, Doshomachi 3-chome, Chuo-ku, Osaka-shi, Osaka 541 (JP).

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- (74) Agent: SEKI, Hideo; Fujisawa Pharmaceutical Co., Ltd., Osaka Factory, 1-6, Kashima 2-chome, Yodogawa-ku, Osaka-shi, Osaka 532 (JP).

### (54) Title: 4,5-DIARYL OXAZOLE DERIVATIVES

#### (57) Abstract

Heterocyclic compounds of formula (I), wherein  $R^1$  is carboxy or protected carboxy,  $R^2$  is aryl which may have suitable substituent(s),  $R^3$  is aryl which may have suitable substituent(s),  $R^4$  is hydrogen, lower alkyl, hydroxy or aryl,  $A^1$  is lower alkylene, (a) is (b), etc.,  $-A^3$ - is (c), etc. and n is 0 or 1, and pharmaceutically acceptable salts thereof which are useful as a medicament.

$$R^{1}-A^{1}-O_{n} \xrightarrow{R^{2}} A^{3} \xrightarrow{N} R^{2}$$
(I)